CO₂ Hydrogenation for Ethanol Production: A Thermodynamic Analysis

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Abstract: The thermodynamics of CO₂ hydrogenation to ethanol synthesis was analyzed by using the principle of Gibbs free energy minimization. According to the reaction mechanism, the product components of the reaction system were determined. The effects of reaction temperature, pressure and the molar ratio of hydrogen to carbon on the equilibrium products were investigated. The results show that methane has a high selectivity in equilibrium products. In order to analyze the influence of reaction conditions on the target product of ethanol, the thermodynamics of CO₂ hydrogenation was studied in methane free products. Since the process of CO₂ hydrogenation is accompanied by the CO hydrogenation reaction (FT synthesis), the CO hydrogenation process was also analyzed and compared with CO₂ hydrogenation. The results show that the CO hydrogenation has more advantages than the CO₂ hydrogenation, and that low temperature and high pressure can improve CO₂/CO conversion and the selectivity of ethanol. The suitable H₂/CO₂ molar ratio in the CO₂ hydrogenation is 3.0-5.0, while the suitable H₂/CO molar ratio in the CO hydrogenation is 0.5-2.0. The comparison of the simulation results with the related experimental results shows that the hydrogenation catalyst needs to be developed continuously to improve the conversion of raw materials and the selectivity to the target product.

Keywords: Thermodynamics, CO₂ Hydrogenation, CO Hydrogenation, Ethanol, Methanol, Dimethyl Ether

1. Introduction

The conversion of carbon dioxide into chemicals and fuels is an important way to reduce climate change and ocean acidification because CO₂ is an important greenhouse gas. In the process of CO₂ resource utilization, CO₂ catalytic hydrogenation is an important field of current C1 chemistry [1, 2]. CO₂ hydrogenation can produce a variety of products such as carbon monoxide, carboxylic acids, aldehydes, alcohols, and hydrocarbons [3-5]. The hydrogenation of CO₂ to methane technology is more mature. CO₂ hydrogenation to methanol, ethanol and dimethyl ether fuel is a promising technology. Methanol synthesis by CO₂ hydrogenation is an effective way to transform CO₂ in industry [6-10]. According to the reaction mechanism, the produced methanol can be further dehydrated to produce dimethyl ether, which can break the thermodynamic equilibrium of methanol synthesis and inhibit the side reaction through the produced water [11-14].

Ethanol is a kind of high quality power fuel, and the

development of fuel ethanol technology is one of the measures to reduce environmental pollution. At present, the methods of ethanol production are mainly crop fermentation and ethylene hydration method. The fermentation method uses corn and other grain crops as raw materials, which will be subject to food security and rising food prices. For the ethylene hydration method, a large amount of sulfuric acid is produced during ethylene hydration, which can corrode the equipment. Therefore, the development of CO₂/CO hydrogenation to ethanol is of great significance.

The study of catalytic hydrogenation of CO₂ to low alcohols is focused on the development of catalysts and thermodynamic studies. Rh based catalysts were used for CO₂ hydrogenation and CO catalytic hydrogenation to ethanol [15-20]. There are few thermodynamic researches on the synthesis of ethanol by CO₂ hydrogenation. An experimental and thermodynamic study for CO₂ conversion to methane and CO using Cu-K/Al₂O₃ catalyst has been

reported [21]. Jia et al [22] carried out the thermodynamics analysis and experimental validation for complicated systems in CO₂ hydrogenation process. The thermodynamic research on CO₂ hydrogenation to low alcohols is based on some reactions of the same type products, considering methanol, ethanol, propanol and C₄ alcohols in the products. However, the actual CO₂ hydrogenation process is complex and produces various products simultaneously. To analyze the real CO₂ hydrogenation process, the suitable hydrogenation process and reaction product components were selected based on the reaction mechanism. The thermodynamic analysis of the influence of reaction conditions on the equilibrium product system was carried out and compared with the experimental results.

2. Calculation Principles and Methods

2.1. Reaction Mechanism Analysis

CO₂ hydrogenation to ethanol passes through the intermediate substance CO ($\text{CO}_2 + \text{H}_2 \leftrightarrow \text{CO} + \text{H}_2\text{O}$) and then continues to be hydrogenated by CO to ethanol. Kusama et al [23] studied the reaction mechanism of CO₂ hydrogenation to ethanol on Rh-Fe/SiO₂ catalyst, as shown in Figure 1.

Figure 1. A possible reaction network of CO₂ hydrogenation [23].

Figure 1 presents a schematic diagram of the reaction network of the formation of CH₄, CO, and ethanol during the hydrogenation of CO₂. In the first step, CO₂ is adsorbed on the catalyst surface by the reverse water gas shift reaction. The adsorbed CO* undergoes successive insertions of H to form an adsorbed CH₃* (methyl) and an adsorbed CH₃CO* (acyl) groups. The hydrogenation of CH₃* produces CH₄ and the acyl group hydrogenation leads to ethanol [15]. The main reactions occurring in the CO₂ hydrogenation system are shown in Table 1.

Table 1. The main reactions in the CO₂ hydrogenation system.

No.	Reaction Equation	$\Delta G_{298\text{ K}}$ (KJ/mol)	$\Delta H_{298\text{ K}}$ (KJ/mol)	$K_{298\text{ K}}$
(1)	$\text{CO}_2 + \text{H}_2 \leftrightarrow \text{CO} + \text{H}_2\text{O}$	28.6	41.1	9.67×10^{-6}
(2)	$\text{CO} + 3\text{H}_2 \leftrightarrow \text{CH}_4 + \text{H}_2\text{O}$	-141.9	-206.0	
(3)	$\text{CO}_2 + 4\text{H}_2 \leftrightarrow \text{CH}_4 + 2\text{H}_2\text{O}$	-113.5	-165.0	7.79×10^{19}
(4)	$\text{CO} + 2\text{H}_2 \leftrightarrow \text{CH}_3\text{OH}$		-90.4	
(5)	$\text{CO}_2 + 3\text{H}_2 \leftrightarrow \text{CH}_3\text{OH} + \text{H}_2\text{O}$	3.5	-49.3	2.45×10^{-1}
(6)	$2\text{CH}_3\text{OH} \leftrightarrow \text{CH}_3\text{OCH}_3 + \text{H}_2\text{O}$		-24.52	
(7)	$\text{CH}_3\text{OCH}_3 + \text{CO} \leftrightarrow \text{CH}_3\text{COOCH}_3$			
(8)	$\text{CH}_3\text{COOCH}_3 + 2\text{H}_2 \leftrightarrow \text{C}_2\text{H}_5\text{OH} + \text{CH}_3\text{OH}$			
(9)	$\text{CH}_3\text{OH} + \text{CO} + 2\text{H}_2 \leftrightarrow \text{C}_2\text{H}_5\text{OH} + \text{H}_2\text{O}$	-97.0	-165.1	
(10)	$2\text{CH}_4 + \text{H}_2\text{O} \leftrightarrow \text{C}_2\text{H}_5\text{OH} + 2\text{H}_2$			
(11)	$\text{C}_2\text{H}_5\text{OH} \rightarrow \text{CH}_3\text{CHO} + \text{H}_2 \leftrightarrow \text{CH}_4 + \text{CO} + \text{H}_2$			
(12)	$2\text{CO} + 4\text{H}_2 \leftrightarrow \text{C}_2\text{H}_5\text{OH} + \text{H}_2\text{O}$	-221.1	-253.6	
(13)	$2\text{CO}_2 + 6\text{H}_2 \leftrightarrow \text{C}_2\text{H}_5\text{OH} + 3\text{H}_2\text{O}$	-32.4	-86.7	4.70×10^5
(14)	$\text{C}_2\text{H}_5\text{OH} + 3\text{H}_2\text{O} \leftrightarrow \text{CH}_3\text{COOH} + 2\text{H}_2$			
(15)	$2\text{CO}_2 + 7\text{H}_2 \leftrightarrow \text{C}_2\text{H}_6 + 4\text{H}_2\text{O}$	-78.7	-132.1	6.26×10^{13}
(16)	$3\text{CO}_2 + 10\text{H}_2 \leftrightarrow \text{C}_3\text{H}_8 + 6\text{H}_2\text{O}$	-70.9	-125.0	2.64×10^{12}
(17)	$\text{CH}_4 \leftrightarrow \text{C} + 2\text{H}_2$		74.9	
(18)	$2\text{CO} \leftrightarrow \text{CO}_2 + \text{C}$		-172.5	

F-T synthesis yields alkanes (C_nH_{2n+2}), olefins (C_nH_{2n}), alcohols (C_nH_{2n+1}OH), aldehydes (C_nH_{2n+1}COH) and carbon deposits (equation 18, 19). In the process of CO₂/CO hydrogenation, methanation is easy to occur. Too high methane is not conducive to the synthesis of ethanol. In order to discuss the influence of reaction conditions on the target product system, the synthesis processes of CO₂ hydrogenation in the products containing methane and in the absence of methane were studied.

Since the process of CO₂ hydrogenation to ethanol is usually accompanied by the processes of CO hydrogenation and the hydrogenation of CO and dimethyl ether (DME) to ethanol synthesis, the following calculations were determined.

(a) CO₂/(CO+DME) hydrogenation to ethanol products containing methane; (b) CO₂/CO hydrogenation to ethanol products in the absence of methane.

2.2. Calculation Methods and Parameters

The Gibbs free energy minimization method is widely used to analyze the equilibrium composition of complex reaction systems. RGibbs reactor model was used by Aspen Plus software in this analysis. In thermodynamic analysis, it is assumed that all the equilibrium components are gas and do not take into account the influence of carbon deposition, implying that the reactions (17, 18) are not considered in Table 1. Considering the existence of nonideal gas in the

system, the Peng-Robinson equation of state is chosen.

In this simulation, the raw material conversion rate and the product selectivity were defined as follows, respectively.

$$X(\text{CO}_X) = [(F_{\text{CO}_X, \text{in}} - F_{\text{CO}_X, \text{out}}) / F_{\text{CO}_X, \text{in}}] \times 100\% \quad (1)$$

$$X(\text{H}_2) = [(F_{\text{H}_2, \text{in}} - F_{\text{H}_2, \text{out}}) / F_{\text{H}_2, \text{in}}] \times 100\% \quad (2)$$

$$S_i = [\lambda F_{i, \text{out}} / (F_{C, \text{in}} - F_{C, \text{out}})] \times 100\% \quad (3)$$

$$S_{\text{H}_2\text{O}} = [2F_{\text{H}_2\text{O}} / (F_{\text{H}_2, \text{in}} - F_{\text{H}_2, \text{out}})] \times 100\% \quad (4)$$

Among which, $X(\text{CO}_X)$ and $X(\text{H}_2)$ are the conversion rates of CO₂/CO and H₂, respectively. F_i is the molar flow rate of species i and the subscripts of in and out respectively indicate the state of being in and out. S_i is the selectivity of carbonaceous substances, in which λ is the number of carbon atoms in species i . $S_{\text{H}_2\text{O}}$ is the selectivity of H₂O calculated according to transformed H₂.

The equilibrium product yield can be defined by the equation of $Y_i = X(\text{CO}_X) \cdot S_i$, indicating that Y_i can be determined by the rate of raw material conversion and the selectivity of the product [24]. Therefore, the effects of reaction conditions on the conversion of raw materials and the selectivity of products were discussed.

3. Results and Discussion

3.1. CO₂/(CO+DME) Hydrogenation to Ethanol Products Containing Methane

For the processes of CO₂ hydrogenation and CO+DME hydrogenation to synthesize ethanol, according to the reaction mechanism in Figure 1 and the equation (1-13) in Table 1, the determined components and their codes of the reaction system were respectively as follows: CO₂, CO, H₂, H₂O, CH₃OH (MeOH), CH₃OCH₃ (DME), C₂H₅OH (EtOH), CH₃COOCH₃ (MA), CH₃CHO (AH) and CH₄.

Figure 2 shows the effects of reaction conditions on CO₂/(CO+DME) hydrogenated products containing methane. As can be seen from Figure 2, for CO₂ hydrogenation and CO+DME hydrogenation, the amount of CH₄ in the equilibrium composition is relatively large. Under the reaction conditions, the selectivity of CH₄ is close to 100%, while the amount of other products is relatively low, which suggests that the reaction is largely thermodynamically beneficial to methanation [25]. For the reaction (3) of (CO₂ + 4H₂ ↔ CH₄ + 2H₂O), the reaction is exothermic and the equilibrium constant is in the order of 10¹⁹ in standard condition. At other temperatures, the equation (3) also has high reaction equilibrium constants.

Figure 2. Effects of reaction conditions on CO₂/(CO+DME) hydrogenated products containing methane. (a, b, c) CO₂ hydrogenation; (d) CO+DME hydrogenation. Reaction conditions: (a) 6.0 MPa, H₂/CO₂ = 3.0; (b) 693 K, H₂/CO₂ = 3.0; (c) 693 K, 7.0 MPa; (d) 6.0 MPa, H₂:CO:DME = 2:1:1.

As shown in Figure 2(a), for CO_2 hydrogenation, with the increase of the reaction temperature, the equilibrium conversion rate of carbon dioxide and hydrogen decreases, and the selectivity of CO increases. The amount of other oxygenated compounds is very low, but their selectivities increase with increasing reaction temperature, and the selectivity of methanol is slightly higher than that of ethanol. The amount of other oxygenated compounds is small, but their selectivity increases with reaction temperature.

As shown in Figure 2(b), increasing the reaction pressure can increase the conversion rate of raw materials and the selectivity of products. It is shown that the total reaction is a reaction of reduced pressure, which can also be seen from several major reactions (3, 5, 13) in Table 1. Increasing the pressure is beneficial to hydrogenation. However, the reaction pressure will affect the equipment investment, production cost and safety, so the working pressure should be properly chosen.

As illustrated in Figure 2(c), feed composition influences its conversion rate and the product selectivity. In the CO_2 hydrogenation process, the CO_2 conversion rate increases with the increase of H_2/CO_2 . After H_2/CO_2 to 4.0, the conversion rate of CO_2 can reach 100%, and the H_2 conversion rate and CO selectivity decrease obviously due to hydrogen excess. In H_2/CO_2 range of 0-10, the selectivity of methane is close to 100%, the selectivity of H_2O is also high, and the selectivity of other products is very low. The appropriate H_2/CO_2 ratio is 3.0-5.0.

As shown in Figure 2(d), in the $\text{CO}+\text{DME}$ hydrogenation process, the conversion rate of CO is close to 100%, but the conversion of DME and H_2 decrease with the increase of temperature. Under the reaction conditions, the selectivities of methane, water and CO_2 are high and the selectivity of other

substances is very low, but they also show a tendency to increase with the increase of the reaction temperature. The effects of pressure and the molar ratio of hydrogen to carbon on the equilibrium product of $\text{CO}+\text{DME}$ hydrogenation are similar to that of CO_2 hydrogenation (Figures omitted). The results of Figure 2(a, d) show that the hydrogenation of CO_2 and $\text{CO}+\text{DME}$ hydrogenation are exothermic reactions, and the water gas shift and its reverse reactions are dominant in the reaction. To improve the conversion rate of raw materials, the reaction should be carried out at low temperature.

3.2. CO_2/CO Hydrogenated Products in the Absence of Methane

In order to analyze the influence of reaction conditions on the target products of CO_2 hydrogenation and CO hydrogenation in detail, the thermodynamic analysis of the processes of CO_2 hydrogenation and CO hydrogenation to ethanol was carried out without considering methane in products. According to the reaction mechanism of Figure 1 and the reaction (4-9) and (12, 13) in Table 1, the material compositions of the reaction system were selected: CO_2 , CO, H_2 , H_2O , CH_3OH (MeOH), CH_3OCH_3 (DME), $\text{C}_2\text{H}_5\text{OH}$ (EtOH) and $\text{CH}_3\text{COOCH}_3$ (MA). The calculated conditions are as follows: the reaction temperature is 293-873 K, the reaction pressure is 0.1-9.0 MPa, and the molar ratio of H/C is 0.5-10.

3.2.1. Effects of Temperature and Pressure

The effects of reaction temperature and pressure on the synthesis equilibrium products in the absence of methane by CO_2/CO hydrogenation are shown in Figure 3.

Figure 3. Effects of temperature and pressure on the synthesis equilibrium products in the absence of methane by CO₂ hydrogenation (a, b, c) and CO hydrogenation (a1, b1, c1). Reaction condition: (a) 3.0 MPa, $H_2/CO_2=4.0$; (a1) 3.0 MPa, $H_2/CO=4.0$; (b, c) $H_2/CO_2=4.0$; (b1, c1) $H_2/CO=4.0$.

As shown in Figure 3(a, b, c), for CO₂ hydrogenation, the equilibrium transformation of CO₂ decreases first and then increases with the increase of the reaction temperature. At a certain temperature, increasing the pressure is beneficial to increase the conversion rate of CO₂. In Figure 3(a), the conversion of hydrogen decreased from 75% to a minimum value of 15% at about 750 K, and then slightly increased. The selectivity of CO increases with the increase of temperature. As shown in Figure 3(a, c), the selectivity of ethanol is close to 100% at low temperature. The selectivity of ethanol decreases with the increase of temperature. Increasing the pressure is beneficial to the temperature range of high ethanol selectivity. The selectivity of ethanol is higher than that of methanol in thermodynamics [22], which can be explained by the equilibrium constant of ethanol synthesis from CO₂ hydrogenation in Table 1 above the corresponding value of methanol synthesis. The selectivities of CH₃OH, CH₃OCH₃ and CH₃COOCH₃ are low, and their selectivities increases first and then decreases with the increase of temperature.

The main reasons for the above trend is that the synthesis of methanol, ethanol and dimethyl ether are all exothermic reactions and the reverse water gas shift reaction (RWGS) is an endothermic reaction. The hydrogenation reaction is dominant at low temperature and the reverse water gas shift reaction is not dominant, showing the conversion rate of CO₂ decrease with the increase of temperature. When the temperature rises to a certain extent, the reaction rate of

reverse water gas shift will be higher than that of hydrogenation reaction, the CO₂ conversion rate is increased with the increase of temperature [22]. Therefore, according to this thermodynamic result, CO₂ hydrogenation synthesis is not suitable for under high temperature conditions, and this requires that industrial hydrogenation catalyst must have good low temperature activity.

As shown in Figure 3(a1, b1, c1), the reaction rule of CO hydrogenation is slightly different from that of CO₂ hydrogenation. In Figure 4(a1), the CO conversion rate is close to 100% at low temperature. When the temperature is above 650 K, the conversion rate of CO decreases rapidly with the increase of temperature, and the conversion rate of hydrogen shows a similar trend. The selectivities of ethanol and water decrease slightly with the increase of temperature. The reason for this phenomenon is that the main reaction is controlled by water vapor shift reaction at high temperature.

As illustrated in Figure 3(b1, c1), Increasing the pressure is beneficial to increase the conversion rate of CO and the selectivity of ethanol. However, when the pressure is higher than 5.0 MPa, the increase of the pressure can not significantly increase the conversion of CO and the selectivity of ethanol.

3.2.2. Effect of Raw Material Composition

The effect of raw material composition on the synthesis of ethanol balanced products in the absence of methane by CO₂/CO hydrogenation is shown in Figure 4.

Figure 4. Effect of raw material composition on the synthesis equilibrium products in the absence of methane by CO₂ hydrogenation (a), CO hydrogenation (b) and (CO₂+CO) hydrogenation (c, d). Reaction condition: 5.0 MPa, 473 K, in addition, (c) H₂/(CO+CO₂)=2.0, (d) H₂/(CO+CO₂)=4.0.

As shown in Figure 4(a, b), the material composition has a great influence on the CO₂/CO hydrogenation process. Under the reaction conditions of 5.0 MPa and 473 K, the conversion rate of CO₂ increases with the change of H₂/CO₂ from 0 to 4.0, and when H₂/CO₂ is above 4.0, the conversion rate of CO₂ almost reach 100% in thermodynamics. CO hydrogenation also showed a similar pattern.

As illustrated in Figure 4(c), under a certain condition of hydrogen to carbon ratio, both CO and H₂ have high conversion rates with the increase of CO content in carbon source. However, the conversion rate of CO₂ decreases with

the increase of CO content in the raw material, which shows that CO is easier to convert than CO₂.

As shown in Figure 4(d), in the case of hydrogen to carbon ratio of 4.0, due to the relatively sufficient hydrogen content, CO and CO₂ have high conversion rates with the increase of CO content, that is, they are basically transformed in thermodynamics. But the conversion rate of H₂ is low, which is mainly due to the occurrence of water gas shift reaction. The amount of CO₂ is completely transformed, but the H₂ conversion rate decreases due to the excess hydrogen.

Table 2. Comparison of CO₂/CO hydrogenation calculation results with experimental results.

Experimental results					Calculation results				
References	Catalyst	Reaction conditions	Conversion rates (%)	Selectivities (%)	Calculation conditions	Conversion rates (%)	Selectivities (%)		
(1) CO ₂ hydrogenation									
[15]	2%Rh -2.5%Fe/TiO ₂	543 K, 1.0 MPa, H ₂ /CO ₂ =1.0, 0.15 g _{cat} /h WHSV =8000 cm ³ /(g _{cat} h)	X(CO ₂) 6.48	CH ₄ 58.7 C ₂ H ₆ +C ₃ H ₈ +C ₄ H ₁₀ 5.88 AH, CO 0.44, 29.3 MeOH, EtOH 1.28, 4.37	533 K, 1.0 MPa, H ₂ /CO ₂ =3.0	X(CO ₂) 74.6	CH ₄ 100		
[15]	2%Rh/TiO ₂	543 K, 2.0 MPa, H ₂ /CO ₂ =1.0, 0.15 g _{cat} /h WHSV =8000 cm ³ /(g _{cat} h)	X(CO ₂) 19.2	CH ₄ 93.3 C ₂ H ₆ +C ₃ H ₈ +C ₄ H ₁₀ 5.18 AH, CO 0, 4.16 MeOH, EtOH 0, 0	533 K, 3.0 MPa, H ₂ /CO ₂ =3.0	X(CO ₂) 74.8	CH ₄ 100		
[16]	5 wt%Rh -Co-Na (1:1:0.5)/SiO ₂	533 K, 5.0 MPa, H ₂ /CO ₂ =3.0, Flow rate =100cm ³ /min	X(CO ₂) 19.8	CH ₄ 36.7 C ₂ H ₆ +C ₃ H ₈ 14.2 MeOH, EtOH 2.0, 8.7 CO 27.1	533 K, 5.0 MPa, H ₂ /CO ₂ =4.0 (CH ₄ free)	X(CO ₂) 96.4	MeOH 0.03 EtOH 99.9 MA 0.02 CO 0.05 DME 0		
[16]	5 wt% Rh-Co (1:1)/SiO ₂	533 K, 5.0 MPa, H ₂ /CO ₂ =3.0, Flow rate =100 cm ³ /min	X(CO ₂) 25.2	CH ₄ 73.0 C ₂ H ₆ +C ₃ H ₈ 2.5 MeOH, EtOH 13.4, 1.2 CO 9.8	533 K, 6.0 MPa, H ₂ /CO ₂ =3.0	X(CO ₂) 74.9	CH ₄ 100 CO 0.002		
[17]	Rh ₁₀ Se/TiO ₂	523 K, 0.047 MPa, H ₂ /CO ₂ =2.0		EtOH 83.0	533 K, 0.1 MPa, H ₂ /CO ₂ =4.0 (CH ₄ free)	X(CO ₂) 27.0	MeOH 0.024 EtOH 66.6 MA 0.03 DME 0 CO 33.3		
(2) CO hydrogenation									
[18]	RMLF/CNTs	593 K, 3.0 MPa, H ₂ /CO=2.0	X(CO) 5.16	CH ₄ 14.6 C ₂ + hydrocarbons 12.5 C ₂ + oxygenates 52.4 MeOH, CO ₂ 1.5, 19.0	573 K, 3.0 MPa, H ₂ /CO=3.0	X(CO) 74.6	CH ₄ 99.99 CO 0.01		
[19]	2%Rh -2%Ce/SiO ₂	453 K, 0.1 MPa, H ₂ /CO=1:2, 0.2 g _{cat} /h, 8 h 30 cm ³ /min,	Activity (μmol/ (g·h))	CO amount converte d=20.9	543 K, 0.1 MPa, H ₂ /CO ₂ =1.0 (CH ₄ free)	X(CO) 91.85	MeOH 0.01 EtOH 74.5 MA 0.04 DME 0 CO ₂ 25.5		

By comparing CO₂ hydrogenation with CO hydrogenation, it can be concluded that CO hydrogenation has advantages over CO₂ hydrogenation [24]. CO hydrogenation have high conversion rate and wider temperature range with higher ethanol selectivity. The thermodynamic optimization conditions are as follows. For the CO hydrogenation reaction, the temperature is below 650 K, the pressure is 1.0-5.0 MPa, and the H₂/CO molar ratio is 0.5-2.0. For the CO₂ hydrogenation, the temperature is below 500 K, the pressure is 1.0-5.0 MPa, and the H₂/CO₂ molar ratio is 3.0-5.0.

3.3. Results Comparison

Table 2 gives the comparison of the results of this calculation with the experimental results. As shown in Table 2, for the establishment of products in hydrogenation reaction system, the results obtained in this study are basically in agreement with the experimental results of catalytic hydrogenation [15-19]. It can be seen that CO₂ and CO hydrogenation have a higher thermodynamic conversion than that of the catalytic experimental reaction under close reaction conditions. The selectivity of the product is closely related to the performance of the catalyst, and the actual hydrogenation reaction is more complicated. At the same time, it is concluded that the performance of CO₂ hydrogenation catalyst needs further improvement to improve the conversion of CO₂/CO and the selectivity of catalyst to the target product.

4. Conclusions

In this paper, according to the reaction mechanism, the thermodynamics of ethanol production by CO₂ hydrogenation was simulated. Since the process of CO₂ hydrogenation contain the processes of CO hydrogenation and CO+DME hydrogenation, different hydrogenation processes are discussed in this paper, in which CO₂ hydrogenation synthesis processes in the products containing methane and in methane free products were emphatically analyzed.

The influences of reaction temperature, pressure and composition of raw material, such as H₂/CO₂ molar ratio, on reaction equilibrium system were analyzed. The change rules of material equilibrium conversion rate and product selectivity with reaction conditions were obtained. The main conclusions of this paper are as follows.

(1) For the CO₂ hydrogenation to ethanol synthesis containing methane, with the increase of the reaction temperature, the equilibrium conversion rate of carbon dioxide and hydrogen decrease, and the selectivity of CO increase. Under the reaction conditions, the selectivity of CH₄ is close to 100%, and water has higher selectivity, while other oxygenated hydrocarbons are little, suggesting that methanation is thermodynamically easy to occur.

(2) For the CO+DME hydrogenation to ethanol synthesis containing methane, the reaction law similar to that of CO₂ hydrogenation was obtained. The conversions of H₂ and dimethyl ether decrease with the increase of the reaction temperature. Under the reaction conditions, CO has a

conversion rate of 100%, methane has nearly 100% selectivity and the other substances are similar to that of CO₂ hydrogenation.

(3) For the CO₂ hydrogenation to ethanol products in the absence of methane, the equilibrium transformation of CO₂ decreases first and then increases with the increase of the reaction temperature. The selectivity of ethanol is close to 100% at low temperature and then decreases with the increase of temperature. The selectivity of ethanol is higher than that of methanol and Increasing the pressure is beneficial to improve the temperature range with high ethanol selectivity.

(4) The comparison of CO hydrogenation with CO₂ hydrogenation in methane free products, the results show that CO hydrogenation has advantages over CO₂ hydrogenation. CO hydrogenation has a wide range of temperature with a high conversion rate and a high selectivity for ethanol. The thermodynamic optimization conditions are as follows. For the CO₂ hydrogenation reaction, the temperature is below 500 K, the pressure is 1.0-5.0 MPa, and the H₂/CO₂ molar ratio is 3.0-5.0. For the CO hydrogenation reaction, the temperature is below 650 K, the pressure is 1.0-5.0 MPa, and the H₂/CO molar ratio is 0.5-2.0.

(5) The thermodynamic results were compared with the experimental results, which indicating that the hydrogenation catalysts need to be improved to increase the conversion of the raw materials and the selectivity of the target product.

(6) The reaction system of CO₂ hydrogenation is complex and produces many kinds of products. Carbon deposition is easy to occur at low temperature, which can lead to deactivation of catalyst. Based on the more accurate catalyst reaction mechanism and considering carbon deposition in the product system, the thermodynamic analysis and kinetic study should be further developed.

Nomenclature

MeOH	methanol
DME	dimethyl ether
EtOH	ethanol
AH	acetaldehyde
H ₂ /CO ₂	the molar ratio of H ₂ /CO ₂
RWGS	reverse water gas shift reaction
$F_{i,in}$	molar flow rate of the <i>i</i> th species in feed (mol/s)
$F_{i,out}$	molar flow rate of the <i>i</i> th species in products (mol/s)
F_C	molar flow rate of carbon (mol/s)
$\Delta G_{298\text{ K}}$	standard Gibbs free energy change (kJ/mol)
$\Delta H_{298\text{ K}}$	standard enthalpy change (kJ/mol)
$K_{298\text{ K}}$	standard equilibrium constant
P	pressure (MPa)
S	selectivity of the product
T	temperature (K)
X	conversion rate
$X(\text{H}_2)$	H ₂ conversion rate
Y_i	equilibrium yield of product
λ	the number of carbon atoms in the <i>i</i> th species in products

References

- [1] Omae, Iwao. "Recent developments in carbon dioxide utilization for the production of organic chemicals". *Coord. Chem. Rev.*, Vol. 256, No. 13–14, 2012, pp. 1384-1405.
- [2] Sternberg, A., C. Jens, and A. Bardow. "Life Cycle Assessment of C1 Chemicals from Hydrogen and Carbon Dioxide". *Chemie Ingenieur Technik*, Vol. 88, No. 9, 2016, pp. 1343-1344.
- [3] Wang S, Mao D S, Guo X M, & Lu G Z. "Dimethyl ether synthesis from CO₂ hydrogenation over CuO-TiO₂-ZrO₂/HZSM-5 catalysts". *Acta Physico-Chimica Sinica*, Vol. 27, No. 11, 2011, pp. 2651-2658.
- [4] Swapnesh, Anang, V. C. Srivastava, and I. D. Mall. "Comparative Study on Thermodynamic Analysis of CO₂ Utilization Reactions". *Chem. Eng. Technol.*, Vol. 37, No. 10, 2015, pp. 1765-1777.
- [5] G. Zahedi, A. Elkamel, and A. Lohiș. "Dynamic Optimization Strategies of a Heterogeneous Reactor for CO₂ Conversion to Methanol". *Energy & Fuels*, Vol. 21, No. 5, 2007, pp. 2977-2983.
- [6] Jia C, Gao J, Dai Y, Zhang J, & Yang Y. "The thermodynamics analysis and experimental validation for complicated systems in CO₂ hydrogenation process". *Journal of Energy Chemistry*, Vol. 25, No. 6, 2016, pp. 1027-1037.
- [7] Matej Huš, Venkata D B C. Dasireddy, Neja Strah Štefančič, & Blaž Likozar. "Mechanism, kinetics and thermodynamics of carbon dioxide hydrogenation to methanol on Cu/ZnAl₂O₄ spinel-type heterogeneous catalysts". *Applied Catalysis B: environmental*, Vol. 207, 2017, pp. 267-278.
- [8] Toyir J, Piscina P R D L, Fierro J L G, & Homs N. "Catalytic performance for CO₂ conversion to methanol of gallium-promoted copper-based catalysts: influence of metallic precursors". *Applied Catalysis B Environmental*, Vol. 34, No. 4, 2001, pp. 255-266.
- [9] Wu J, Saito M, Takeuchi M, & Watanabe T. "The stability of Cu/ZnO-based catalysts in methanol synthesis from a CO₂-rich feed and from a CO-rich feed". *Applied Catalysis A General*, Vol. 218, No. 1, 2001, pp 235-240.
- [10] Liu XM, Lu GQ, Yan ZF, Beltramini J. "Recent Advances in Catalysts for Methanol Synthesis via Hydrogenation of CO and CO₂". *Ind. Eng. Chem. Res.*, Vol. 42, No. 25, 2003, pp. 6518-6530.
- [11] Fu Y, Hong T, Chen J, Auroux A, & Shen J. "Surface acidity and the dehydration of methanol to dimethyl ether". *Thermochimica Acta*, Vol. 434, No. 1-2, 2005, pp. 22-26.
- [12] Zhang L, Zhang H, Ying W, & Fang D. "Dehydration of methanol to dimethyl ether over γ -Al₂O₃ catalyst: intrinsic kinetics and effectiveness factor". *Can. J. Chem. Eng.*, Vol. 91, No. 91, 2013, pp. 1538–1546.
- [13] Kochkin, Y. N., Vlasenko, N. V., Struzhko, V. L., Puziy, A. M., & Strizhak, P. E. "Methanol carboxylation over zirconium dioxide: effect of catalyst phase composition on its acid - base spectrum and direction of catalytic transformations". *Can. J. Chem. Eng.*, Vol. 94, No. 4, 2016, pp. 745–751.
- [14] Ghodhbene M, Bougie F, Fongarland P, & Iliuta M C. "Hydrophilic zeolite sorbents for in - situ water removal in high temperature processes". *Can. J. Chem. Eng.*, Vol. 95, No. 10, pp 1842–1849, 2017.
- [15] Makarand R. Gogate, Robert J. Davis. "Comparative study of CO and CO₂ hydrogenation over supported Rh-Fe catalysts". *Catal. Commun.*, Vol. 11, 2010, pp. 901–906.
- [16] Kusama, Hitoshi, and H. Arakawa. "Hydrogenation of CO₂ over SiO₂ Supported Rh-Co-alkalimetal Catalysts." *Nippon Kagaku Kaishi*, Vol. 2002, No. 1, 2002, pp. 107-110.
- [17] Izumi Y. "Selective Ethanol Synthesis from Carbon Dioxide: Roles of Rhodium Catalytic Sites". *Platinum Metals Review*, Vol. 41, No. 4, 1997, pp. 166-170.
- [18] Fan Z, Chen W, Pan X, & Bao X. "Catalytic conversion of syngas into C2 oxygenates over Rh-based catalysts—Effect of carbon supports". *Catalysis Today*, Vol. 147, No. 2, 2009, pp. 86-93.
- [19] Wang Yaquan. Study on the synthesis of ethanol through CO hydrogenation catalyzed by Rh based catalysts [J]. *Chinese Journal of Catalysis*, Vol. 20, No. 1, 1999, pp. 7-11. (in chinese).
- [20] Kurakata H, Izumi Y, Aika K. "Ethanol synthesis from carbon dioxide on TiO₂-supported [Rh10Se] catalyst", *Chem. Commun.*, Vol. 3, No. 3, 1996, pp. 389-390.
- [21] Ahmad W, Al-Matar A, Shawabkeh R, & Rana A. "An experimental and thermodynamic study for conversion of CO₂ to CO and methane over Cu-K/Al₂O₃", *J. Environ. Chem. Eng.*, Vol. 4, No. 3, 2016, pp. 2725-2735.
- [22] Jia C, Gao J, Dai Y, Zhang J, & Yang Y. "The thermodynamics analysis and experimental validation for complicated systems in CO₂ hydrogenation process". *Journal of Energy Chemistry*, Vol. 25, No. 6, 2016, pp. 1027-1037.
- [23] Kusama H, Okabe K, Sayama K, Arakawa H. "Ethanol synthesis by catalytic hydrogenation of CO₂ over Rh-Fe/SiO₂ catalysts". *Energy*, Vol. 22, 1997, pp. 343–348.
- [24] Guangxin Jia, Yisheng Tan, Yizhuo Han. "A comparative study on the Thermodynamics of dimethyl ether synthesis from CO Hydrogenation and CO₂ Hydrogenation". *Ind. Eng. Chem. Res.*, Vol. 45, 2006, pp. 1152-1159.
- [25] Torrente-Murciano L, Mattia D, Jones M D, & Plucinski P K. "Formation of hydrocarbons via CO₂ hydrogenation – A thermodynamic study". *Journal of CO₂ Utilization*, Vol. 6, No. 3, 2014, pp. 34-39.